Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claims 1 - 58 (Cancelled)

59. (Currently Amended) A method of reducing risk of, preventing or treating, non-proliferative diabetic retinopathy and/or macular edema, in a mammal by administrating an effective amount of a medicament comprising at least one compound capable of inhibiting the visual cycle, to said mammal, wherein the at least one compound comprises a retinoid of the formula V:

wherein R1, R4, R9, R10 and R12 is CH3, and R3, R5, R6, R7, and R8 is H, and

wherein R11 is selected from the group consisting of:

- -COOH,
- an alcohol group,
- -CHO,
- -CH2OCOCH2Br,
- -CH2OCOCH2Cl,
- -COOCH2CH3,
- -CONHR', wherein R' is 4-hydroxy-phenyl or ethyl, and

-COOR", wherein R" is beta-D-glucuronide, and wherein the configuration of the four isoprenoid units is all trans (E) or one or more is cis (Z), with the proviso that when R11 is -COOH, the configuration is not 9 cis (2E, 4E, 6Z 8E) or all trans.

- 60. (Previously Presented) The method of claim 59, wherein said mammal is a human being.
- 61. (Previously Presented) The method of claim 59, wherein said mammal has been diagnosed with diabetes.

62-78. (Cancelled)

79. (Currently Amended) The method of claim 59, wherein the at least one compound comprises a compound selected from the group consisting of: isotretinoin (13-cis-retinoic acid), 11-cis-retinol, 11-cis-retinal, 11-cis-retinyl bromoacetate, acitretin, etretinate, fenretinide, 4-oxo-isotretinoin, motretinide, retinaldehyde, all-trans-retinyl bromoacetate, all-trans-retinyl chloroacetate, and retinoyl betaglucoronide.

80-107. (Cancelled)

- 108. (Previously Presented) The method of claim 59, wherein the at least one compound is composed as a pro-drug.
- 109. (Previously Presented) The method of claim 59, wherein the medicament is in a form for being administered locally.
- 110. (Previously Presented) The method of claim 109, wherein the medicament is in a form for being administered intravitreally.
- 111. (Previously Presented) The method of claim 59, wherein the medicament is in device formulation held confined by mechanical or physico-chemical effects.

- 112. (Previously Presented) The method of claim 59, wherein the medicament is in a slow-release formulation.
- 113. (Previously Presented) A method comprising a pharmaceutical composition suitable for intravitreal implantation comprising a pharmaceutically effective amount of at least one compound capable of inhibiting the visual cycle and/or dark adaptation.
- 114. (Previously Presented) The method of claim 113, wherein said pharmaceutically effective amount of said at least one compound is determined by measuring the level of reduction of dark adaptation in a treated subject.
- 115. (Previously Presented) The method of claim 113, wherein said pharmaceutical composition is in device formulation held confined by physico-chemical effects.
- 116. (New). The method of claim 59 which is a method of treating non-proliferative diabetic retinopathy and/or macular edema.
- 117. (New). The method of claim 59 which is a method of treating non-proliferative diabetic retinopathy.
- 118. (New). The method of claim 59 wherein the alcohol group is $-\text{CH}_2\text{OH}$.